Detailed and Complete Listing of Claims:

1-3. (Canceled).

4. (Currently Amended) ~~The preparation according to claim 2~~ A percutaneous absorption preparation which comprises a skin contacting base containing a compound having angiotensin II antagonistic activity and a skin permeability regulator, and a support, wherein the skin permeability regulator ~~which~~ comprises a fatty acid ester, a polyol and a nonionic surfactant ~~as the skin permeability regulator~~.

5. (Canceled).

6. (Currently Amended) The preparation according to claim ~~1~~ 4, wherein the compound having angiotensin II antagonistic activity is a non-peptide compound.

7. (Withdrawn-Currently Amended) The preparation according to claim ~~1~~ 4, wherein the compound having angiotensin II antagonistic activity is 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]benzimidazole-7-carboxylic acid or a salt thereof.

8. (Currently Amended) The preparation according to claim ~~1~~ 4, wherein the compound having angiotensin II antagonistic activity is 1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]benzimidazole-7-carboxylate or a salt thereof.

9. (Withdrawn-Currently Amended) The preparation according to claim ~~1~~ 4, wherein the compound having angiotensin II antagonistic activity is 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl]methyl]benzimidazole-7-carboxylic acid or a salt thereof.

10. (Canceled).

11. (Currently Amended) The preparation according to claim ~~10~~ 4, wherein the fatty acid ester is an ester of C_{10-22} carbonic acid and C_{1-12} alkylalcohol.

12. (Currently Amended) The preparation according to claim ~~10~~ 4, wherein the fatty acid ester is isopropyl myristate, isopropyl palmitate, butyl myristate or diethyl sebacate.

13. (Currently Amended) The preparation according to claim ~~10~~ 4, wherein the fatty acid ester is isopropyl myristate.

14. (Canceled).

15. (Currently Amended) The preparation according to claim ~~14~~ 4, wherein the polyol is ethylene glycol, propylene glycol, 1,3-butylene glycol, polyethylene glycol or glycerin.

16. (Currently Amended) The preparation according to claim ~~14~~ 4, wherein the polyol is propylene glycol.

17. (Canceled).

18. (Currently Amended) The preparation according to claim ~~17~~ 4, wherein the nonionic surfactant is a fatty acid amide, a polyol fatty acid ester or a polyglycerol fatty acid ester.

19. (Currently Amended) The preparation according to claim ~~17~~ 4, wherein the nonionic surfactant is a fatty acid amide.

20. (Original) The preparation according to claim 19, wherein the fatty acid amide is lauric acid diethanol amide or a material containing the same.

21. (Original) The preparation according to claim 20, wherein lauric acid diethanol amide or a material containing the same is palm fatty acid diethanol amide.

22. (Currently Amended) The preparation according to claim ~~1~~ 4 which is a skin patch.

23. (Currently Amended) The preparation according to claim ~~10~~ 4, wherein the amount of the fatty acid ester in the skin contacting base is about 1 to 30% by weight based on the weight of the skin contacting base.

24. (Currently Amended) The preparation according to claim ~~14~~ 4, wherein the amount of the polyol in the skin contacting base is about 1 to 30% by weight based on the weight of the skin contacting base.

25. (Currently Amended) The preparation according to claim ~~17~~ 4, wherein the amount of the nonionic surfactant in the skin contacting base is about 1 to 15% by weight based on the weight of the skin contacting base.

26. (Currently Amended) The preparation according to claim ~~1~~ 4 which further contains an adhesive in the skin contacting base.

27. (Original) The preparation according to claim 26, wherein the adhesive is an acrylic adhesive.

28. (Original) The preparation according to claim 26, wherein the adhesive is a self cross-linking acrylic adhesive.

29. (Currently Amended) A preparation according to claim ~~1~~ 4, wherein the amount of the compound having angiotensin II antagonistic activity in the skin contacting base is about 0.01 to 70% by weight based on the weight of the skin contacting base.

30. (Currently Amended) The preparation according to claim ~~1~~ 4, wherein the amount of the skin permeability regulator in the skin contacting base is about 0 to 70% by weight based on the weight of the skin contacting base.

31. (Original) The preparation according to claim 26, wherein the amount of the adhesive in the skin contacting base is about 5 to 99% by weight based on the weight of the skin contacting base.

32. (Currently Amended) The preparation according to claim ~~1~~ 4, wherein the amount of the compound having angiotensin II antagonistic activity per unit of skin contacting area in the skin contacting base is about 0.01 to 100mg/cm^2.

33. (Currently Amended) The preparation according to claim ~~1~~ 4 which maintains effective concentration of the compound having angiotensin II antagonistic activity in blood for one day or more.

34. (Currently Amended) A method of treating angiotensin II-mediated diseases which comprises administrating a percutaneous absorption preparation comprising a skin contacting base containing a compound having angiotensin II antagonistic activity and a skin permeability regulator, and a support, wherein the skin permeability regulator comprises a fatty acid ester, a polyol, and a nonionic surfactant.

35-37. (Canceled).

38. (Currently Amended) A method of percutaneous absorption of a compound having angiotensin II antagonistic activity which comprises adding a compound having angiotensin II antagonistic activity and a skin permeability regulator to a percutaneous absorption preparation comprising a skin contacting base and a support, wherein the skin permeability regulator comprises a fatty acid ester, a polyol, and a nonionic surfactant.

39. (Original) A method of regulating percutaneous absorption of a compound having angiotensin II antagonistic activity, which comprises adding a fatty acid ester, a polyol and a nonionic surfactant to a percutaneous absorption preparation comprising the compound having angiotensin II antagonistic activity.

40. (Canceled).